

Please REPLACE Claim 16 with the following AMENDED Claim 16:

Marked-up copy of amended claim 16:

16. (Amended) A method of screening a patient for cancer, the method comprising:

a) performing an amplification technique on a sample from a biopsy taken from a patient to produce an amplified sample, wherein the sample comprises nucleic acid, and wherein the amplification technique is specific for amplification of a portion of a HPV16 sequence, and at least one HPV sequence selected from the group consisting of HPV18, HPV31, HPV 33, HPV35, HPV45, HPV58,

wherein said amplification technique comprises use of a primer set comprising either SEQ. ID. NOS.:1 and 2 or SEQ. ID. NOS.: 3 and 4.

Clean copy of amended claim 16:

16. A method of screening a patient for cancer, the method comprising:

a) performing an amplification technique on a sample from a biopsy taken from a patient to produce an amplified sample, wherein the sample comprises nucleic acid, and wherein the amplification technique is specific for amplification of a portion of a HPV16 sequence, and at least one HPV sequence selected from the group consisting of HPV18, HPV31, HPV 33, HPV35, HPV45, HPV58,

sequence, and at least one HPV sequence selected from the group consisting of HPV18, HPV31, HPV 33, HPV35, HPV45, HPV58,

wherein said amplification technique comprises use of a primer set comprising either SEQ. ID. NOS.:1 and 2 or SEQ. ID. NOS.: 3 and 4.

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